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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/567,230	02/03/2006	Gordon Bell	70257	9450

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SYNGENTA CROP PROTECTION, INC.  
PATENT AND TRADEMARK DEPARTMENT  
410 SWING ROAD  
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EXAMINER
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PRYOR, ALTON NATHANIEL

ART UNIT	PAPER NUMBER
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1616

MAIL DATE	DELIVERY MODE
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08/27/2008

PAPER

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.



### **DETAILED ACTION**

Applicant's arguments filed 5/12/08 have been fully considered but they are not persuasive. Previous rejections not addressed below are withdrawn.

#### ***Claim Rejections - 35 USC § 102***

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 1-5,9,10,12 remain rejected under 35 U.S.C. 102(b) as being anticipated by Aven (EP 1023832; 8/2/00). Aven teaches an aqueous, concentrate suspension comprising an alkylpolyglycosides (oil based adjuvant) and a hydrotrope. See page 5 paragraph 31. Aven teaches that the concentrate can comprise approximately 25% adjuvant (calculated from the addition of (g/L): 400 active, 500 adjuvant, 100 surfactant, and 800; then divided total value into 500 g/L adjuvant to arrive at about 25% adjuvant). See abstract. A suspension equates to dispersion of active particles (solid) in a liquid which meets the limitation of second phase (solid) being dispersed in a continuous phase. See abstract, paragraph 53. Aven also teaches the addition of a liquid active ingredient to the composition which meets the claim limitation of the second phase comprising a water-immiscible liquid. See paragraph 26. With respect to claim 12 the composition in about 25% oil based adjuvant meets the limitation of the oil base adjuvant comprising a dispersed agrochemical concentrate therein.

*Response to Applicants Arguments*

The Applicants argue that Aven does not motivate a skill person to replace water soluble APGs with an oil based adjuvant. The Examiner argues that alkylpolyglycosides (APGs) such as Glucopons (polymeric structure) are combined with hydrotropes. Glucopons are not water soluble; therefore, Glucopons are structurally considered lipophilic or hydrophobic which would characterize them as being oil soluble or oil-based compounds. See Aven page 5 lines 26-28.

***Claim Rejections - 35 USC § 103***

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 6,13-15 remain rejected under 35 U.S.C. 103(a) as being unpatentable over Aven (EP 1023832) as applied to claims 1-5,9,10 above. Aven teaches that the suspension concentrates are processed by well established procedures including mixing and / or milling of actives with other substances such as solvents and adjuvants. See paragraphs 53,63,64 . Aven teaches that the concentrate can be diluted in water. See paragraph 6. Aven does not specifically teach a) that the continuous phase is prepared first, b) the milling of the solid in water, c) the dilution of the concentrate in a spray tank of water or encapsulation of ingredients (the second phase). The specification does not provide results related to the formulation of the continuous phase first versus the methodology as recited in Aven. In the absence of such results, it is

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obvious that the ordering of the steps will yield the same concentrate (possessing the same chemical and physical characteristics) since Aven and instant invention teaches the mixing / milling of the same chemicals. Both inventions disclose that the concentrate is diluted in water. Therefore whether the concentrate is diluted in spray tank or in some container is immaterial, i.e. the composition should be the same. In the absence of unexpected results, the concentrate diluted in the spray tank should be identical to the concentrate diluted in any other container. With respect to micro-encapsulation (encapsulation), it is standard practice to encapsulate materials to delay their release. This signifies a common practice in the herbicide art. There is nothing unobvious about encapsulating materials in the herbicide art. (e.g. see USPN 5708073).

*Response to Applicants' Argument*

The Applicants argue that Aven does not motivate a skill person to replace water soluble APGs with an oil based adjuvant. The Examiner argues that alkylpolyglycosides (APGs) such as Glucopons (polymeric structure) are combined with hydrotropes. Glucopons are not water soluble; therefore, Glucopons are structurally considered lipophilic or hydrophobic which would characterize them as being oil soluble or oil-based compounds. See Aven page 5 lines 26-28.

***Claim Objection***

Claims 7,8 and 11 remain objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims. The prior does not teach or

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suggest the instant invention comprising the second phase as a micro-emulsion or a third phase.

**THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

***Telephonic Inquiry***

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Alton N. Pryor whose telephone number is 571-272-0621. The examiner can normally be reached on 8:00 a.m. - 4:30 p.m..

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Johann Richter can be reached on 571-272-0646. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Alton N. Pryor/  
Primary Examiner, Art Unit 1616